DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances Application: Akorn, Inc.

[Docket No. DEA-392]

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 7, 2019, Akorn Inc., 1222 West Grand Avenue Decatur, Illinois 62522-1412 applied to be registered as an importer of the following basic class of controlled substance:

Controlled Substance	Drug Code	Schedule
Remifentanil	9739	II

The company plans to import the above listed controlled substance for research purposes.

Dated: August 9, 2019

Neil D. Doherty
Acting Assistant Administrator.
[FR Doc. 2019-18453 Filed: 8/26/2019 8:45 am; Publication Date: 8/27/2019]